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-Figures 3A-and 3B illustrate the HY2 locus of Arabidopsis. Figure 3A shows a map of the region of chromosome 3 containing HY2. Two distinct mapping populations were screened, and mapping results with molecular markers are summarized schematically, indicating that HY2 lies in a region 66 kb in length. Markers starting with the letter c are CAPS markers developed during this study. DNA sequence information for bacterial artificial chromosomes (BACs) MZB10 and F3L24 is available in GenBank/EMBL/DDBJ. The HY2 gene structure with mutations is illustrated at the bottom. Exons are depicted as dark boxes and thick lines, which reflect coding regions and 59/39 untranslated regions, respectively. Dotted lines indicate introns. Figure 3B shows the genomic sequence of HY2 (SEQ ID NO:32) and the deduced HY2 protein sequence (SEQ ID NO:33) from the Columbia (Col) ecotype. Uppercase letters represent exons determined by sequence analysis of HY2 cDNAs. Introns and spacer sequences are indicated with lowercase letters. The stop codon is double underlined. Mutations in hy2 alleles are shown in boldface letters. Single nucleotide polymorphisms in both Ler and Wassilewskija (Ws) ecotypes include the following: inserted T (at nucleotide 234), G364T conversion with amino acid change to Asn, and G1182A conversion (silent). Single nucleotide polymorphisms in the Ler ecotype only include the following: C515A (in intron), G884A (silent), C1145T (in intron), and G1717A (in intron). The single nucleotide polymorphism in Ws ecotype only is C1910T (silent).

Delete the paragraph at page 13, line 25 through page 14, line 13, and insert the following:

Figure 5 shows an alignment of HY2 and HY2-Related Proteins. Alignment of the HY2 protein (SEQ ID NO:34) with proteins of unknown function from oxygenic photosynthetic bacteria identified by PSI BLAST. Conserved residues in 100 or 80% of the aligned sequences are depicted in the consensus sequence with uppercase or lowercase letters, respectively. Sequence similarity groups shown in the consensus sequence reflect conservation in 100% of the sequences. These are labeled as follows: 1, D 5 N; 4, R 5 K; 5, F 5 Y 5 W; and 6, L 5 I 5 V 5 M. Dark shading with white letters, gray shading with white letters, and gray shading with black letters reflect 100, 80, and 60% sequence conservation, respectively. Sequence identifiers correspond to hypothetical proteins from *Synechococcus* sp WH8020 (YCP2\_SYNPY (SEQ ID NO:35) and YCP3\_SYNPY (SEQ ID NO:36)), from *Prochlorococcus* (YHP2\_PROMA (SEQ ID NO:37) and YHP3\_PROMA (SEQ ID NO:38)), and from *Synechocystis* sp PCC 6803 gene (cyanobase locus slr0116 (SEQ ID NO:39); see http://www.kazusa.or.jp/cyano/cyano.html). Database accession numbers are AB045112 for HY2

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al cont (DDBJ), Q02189 for YCP2\_SYNPY (SWISSPROT), Q02190 for YCP3\_SYNPY (SWISSPROT), CAB95700.1 for YHP2\_PROMA (EMBL), CAB95701.1 for YHP3\_PROMA (EMBL), and S76709 for slr0116 (Protein Information Resource). Asterisks are indicated every 20 residues

Delete the paragraph at page 15, lines 1-27, and insert the following:

Figure 10 shows a multiple sequence alignment of the identified bilin reductases. All identified sequences were aligned using the programs CLUSTAL W and MEME. Conserved residues in 90 or 70% of the aligned sequences are depicted in the consensus sequence with uppercase or lowercase letters, respectively. Sequence similarity groups, labeled 1 (D, E), 2 (R, K), 3 (F, Y, W), and 4 (L, I, V, M)., shown in the consensus sequence reflect conservation in >90% of the sequences. Dark shading with white letters, gray shading with white letters, and gray shading with black letters reflect 90, 70, and 50% sequence conservation, respectively. SYNY3, Synechocystis sp PCC6803; SYNPY, Synechococcus sp WH8020; SYN81, Synechococcus sp WH8102; PROMA, Prochloroccocus sp SS120; PROME, Prochloroccocus sp MED4; NOSPU, Nostoc punctiforme; ANASP, Anabaena sp PCC7120; ARATH, Arabidopsis thaliana; and HORVU, Hordeum vulgare. Database accession numbers are GB: AF339056 for PcyA\_ANASP (SEQ ID NO:40) (CyanoBase contig 362), GB: AF339057 for PcyA\_NOSPU (SEQ ID NO:41) (JGI contig 632), PIR: S76709 for PcyA\_SYNY3 (SEQ ID NO:42), PcyA\_SYN81 (SEQ ID NO:43) is on JGI contig 51, GB: AF352050 for PcyA\_PROME (SEQ ID NO:44) (JGI contig 26), SW: Q02189 for PebA\_SYNPY (SEQ ID NO:45), PIR: S31075 (fragment)/ JGI contig 72 for PebA\_SYN81 (SEQ ID NO:46), EMB: CAB95700.1 for PebA\_PROMA (SEQ ID NO:47), PebA\_PROME (SEQ ID NO:48) is on JGI contig 26, GB: AF352049 for PebA\_NOSPU (SEQ ID NO:49) (JGI contig 622), SW: Q02190 for PebB\_SYNPY (SEQ ID NO:50), PebB\_SYN81 (SEQ ID NO:51) is on JGI contig 72, EMB: CAB95701.1 for PebB\_PROMA (SEQ ID NO:52), PebB\_PROME (SEQ ID NO:53) is on JGI contig 26, GB: AF339058 for PebB\_NOSPU (SEQ ID NO:54) (JGI contig 622), DDBJ: AB045112 for HY2\_ARATH (SEQ ID NO:55), EMB: CAB77705.1 for RCCR\_HORVU (SEQ ID NO:56), EMB: CAB16763.1 for RCCR\_ARATH (SEQ ID NO:57). Asterisks indicate every tenth amino acid; dashes indicate gaps; numbers above the line indicate amino acid sequence numbering starting with number one.-

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